Febrivac 3-PLUS suspensija injekcijām ūdelēm

Not authorised

- Mink enteritis virus, Inactivated
- Clostridium botulinum, type C, toxoid
- Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
- Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
- Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated

Product identification

Medicine name:
Febrivac 3-PLUS suspensija injekcijām ūdelēm

Active substance:
Mink enteritis virus, Inactivated

Clostridium botulinum, type C, toxoid

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated

Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated

Target species:
Mink

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Mink enteritis virus, Inactivated
10000.00 50% tissue culture infectious dose / 1.00 unit(s)

Clostridium botulinum, type C, toxoid
0.50 relative unit(s) / 1.00 unit(s)

Pseudomonas aeruginosa, serotype 5, strain EP3, Inactivated
100000000.00 cells / 1.00 unit(s)

Pseudomonas aeruginosa, serotype 6, strain EP2, Inactivated
100000000.00 cells / 1.00 unit(s)

Pseudomonas aeruginosa, serotype 7/8, strain EP1, Inactivated
100000000.00 cells / 1.00 unit(s)

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI20CL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Revoked

Authorised in:

Latvia

Package description:

Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
IDT Biologika GmbH

Marketing authorisation date:
9/03/1999

Manufacturing sites for batch release:
IDT Biologika GmbH

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/99/0944

Date of authorisation status change:
7/05/2024

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.